

**GOODMEDS COUGH RELIEF DEXTROMETHORPHAN HBR- dextromethorphan hydrobromide capsule, gelatin coated
Cabinet Health P.B.C.**

Goodmeds Cough Relief, Dextromethorphan HBr

Drug Facts

Active ingredient (in each softgel)

Dextromethorphan HBr, 15 mg

Purpose

Cough suppressant

Uses

temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold.

Warnings

Do not use

if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- a cough that occurs with too much phlegm (mucus)
- a cough that lasts or is chronic as occurs with smoking, asthma, or emphysema

Stop use and ask a doctor if

cough lasts for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

If pregnant or breast-feeding:

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- do not take more than 8 softgels in any 24-hour period
- this adult product is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and over	take 2 softgels every 6 to 8 hours, as needed
children under 12 years	do not use

Other information

- store at 20-25°C (68-77°F)
- avoid excessive heat above 40°C (104°F)
- protect from light

Inactive ingredients

FD&C blue #1, FD&C red #40, gelatin, glycerin, polyethylene glycol 400, propylene glycol, povidone K30, propyl gallate, purified water, sorbitol sorbitan solution

Questions or comments?

call **1-908-242-6108** Mon-Fri 8:00 AM to 5:00 PM EST

Package Labeling:

Drug Facts (continued)

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goodmeds Compare to the active ingredient in **ROBITUSSIN® Long-Acting CoughGels®**

Cough Relief

Dextromethorphan HBr 15 mg
Cough Suppressant

Non-Narcotic
For Ages 12+

60 Softgels

actual size 

NDC 82725-4110-2

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE

Drug Facts

Active ingredient (in each softgel) **Purpose**

Dextromethorphan HBr, 15 mgCough suppressant

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(Continued on back of the label)

This product is not manufactured or distributed by Haleon, owner of the registered trademark Robitussin® Long-Acting CoughGels®.

Distributed By: ORG 12/25
Cabinet Health P.B.C Made in India
Brooklyn, NY 11222 ITEM# GM-4110-2

Lot No. and Exp. Date: see bottom of bottle



Package Labeling:

Drug Facts (continued)

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(Continued on back of the label)

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ITEM# GM-4110-1

Distributed By:
Cabinet Health P.B.C
Brooklyn, NY 11222
ORG 12/25

Made in India



LOT:

EXP:



Blank boxes for LOT and EXP information.

GOODMEDS COUGH RELIEF DEXTROMETHORPHAN HBR

dextromethorphan hydrobromide capsule, gelatin coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82725-4110	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)		DEXTROMETHORPHAN HYDROBROMIDE	15 mg	
Inactive Ingredients				
Ingredient Name			Strength	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GELATIN (UNII: 2G86QN327L)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
POVIDONE K30 (UNII: U725QWY32X)				
PROPYL GALLATE (UNII: 8D4SNN7V92)				
WATER (UNII: 059QF0KO0R)				
Product Characteristics				
Color	red	Score	no score	
Shape	CAPSULE	Size	8mm	
Flavor		Imprint Code	T	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82725-4110-2	60 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	
2	NDC:82725-4110-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	02/01/2026		

Labeler - Cabinet Health P.B.C. (117102391)

Registrant - Cabinet Health P.B.C. (117102391)

